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APR 1 1 1990

Richard G. Shepherd Project Coordinator Conestoga-Rovers & Associates, Inc. O'Hare Corporate Towers One 10400 W Higgins Rd, Suite 103 Rosemont, IL 60018

Dear Mr. Shepherd:

Enclosed are copies of the sampling results, Quality Assurance data, Chain of Custody forms, and map illustrating the sampling locations for the twelve samples taken during U.S. EPA's and Illinois EPA's February 7, 1990 visit to the Johns-Manville-Waukegan, Illinois Site.

Please contact me at (312) 886-4742 if you have any questions concerning this letter or the enclosures.

Sincerely yours,

Brad Bradley Remedial Project Manager

Enclosures

 ∞ : Kurt Neibergall, IEPA

Eric Meyers, Waukegan

1313 4/7/90

bcc: L. Johnson, 5CS-TUB-03

M. Toney, 5CS-TUB-03

B. Kush, IL/IN #3



Cilent:

Illinois Environmental Protection Agency

Division of Land Pollution Control

2200 Churchill Road

Springfield, Illinois 62706

Contact: Kurt Neibergall, Project Manager

Analysis: Routine/Unit Cost

ASBESTOS BULK SAMPLE EVALUATION

POLARIZED LIGHT MICROSCOPY (PLM) TECHNIQUE

Date Analyzed:

2/9/90

Date Received:

2/9/90

Client Reference: Johns Manville

Maywood

Client P.O.#:

RAI Job #:

2-0366-004.01

Type of Sampl	le: Soil/Debr	is Samples	Sampled b	y: Client			Analyst:	Anita Sumpter-Sturgies
						Fibrous F	orms	
Cllent I.D.	Cllent Code	BAL_#	Sample <u>Description</u>	Sample <u>Treatment</u>	Asbestos Type	<u>%</u>	Non-asbestos Type	Non-fibrous <u>Components</u>
Surface grab sample 50' east of NE corner, misc. pit (15' from east edge of roadway)	X-001	900209-0 3A	Two Components: 1) Brown, fibrous, non-friable 2) Grey, fibrous, friab	1) P, H 2) S, H			1) None Detected 2) Cellulose 1-5%	1) Binder, quartz Total %:55-65 2) Binder, quartz Total %:10-20
East edge of road-80 yd. north of NE corner misc. pit (20' west of MW		900209-04A	Two Components: 1) Black and grey, fibrous, non-friable 2) Grey and brown, fibrous, friable	1) P, H 2) S, H	1) Chrysotile 2) Chrysotile		1) Cellulose TR 2) Cellulose TR	1) Binder Total %:50-60 2) Binder, quartz Total %:85-95

#7)

Client: Illinois Environment	Cllent: Illinois Environmental Reference: Johns Manville				Fibrous I	Forms		
Protection Agency Client I.D. Client Coc	le RAI#	Sample <u>Description</u>	Sample Treatme	Asbesto nt <u>Type</u>	s <u>%</u>	Non-asbestos <u>Type</u>	Non-fibrous Components	
15' East of X-003 roadway-pipe/ siding debris 70 yds. from bend in road to west, -surface grab sample	900209-05 A	Two Components: 1) Brown, fibrous, non-friable 2) Grey and brown, fibrous, friable	1) P, H 2) S, H	1) Chryso 2) Chryso Amosite	ile 2) 10-20	1) Wollastonite 1-5% 2) Cellulose 1-5%	1) Binder, quartz Total %:45-55 2) Binder, quartz Total %:45-55	
East edge of X-004 roadway-sludge / pipe-gravelly 30' north of MW5	900209-06A	Two Components: 1) Black, fibrous, non-friable 2) Grey, fibrous, friable	1) P, H 2) S, H		•	1) None Detected 2) None Detected	1) Binder Total %:20-30 2) Binder Total %:10-20	
Undercut-bank- X-005 indutrial canal (south side) approx. 20 ydwest of new overflow structure	900209-07 A	Grey, fibrous, friable	S, H	Chrysotile	50-60	Cellulose 5-10%	Binder Total %:20-30	
1/2 way down X-006 west bank industrial canal -50" from NW corner (surface grab)	900209-08 A	Three Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable 3) Black, fibrous, gelatinous	1) P, H 2) S, H 3) S, H	•		1) Cellulose TR 2) None Detected 3) Cellulose 60-70%	1) Binder Total %:85-95 2) Binder Total %:10-20 3) Binder Total %:20-30	

Acalyst's Signature Austa Sumplex Sugar Page 2 of 5

Client: Illinois Environmental		Reference:	Johns Manville		orms			
Protect Client 1,D.	ion Agency Client Code	RAL #	Sample Description	Sample Treatmen	Asbestos t Type	<u>%</u>	Non-asbestos Type	Non-fibrous Components
Southeast corner-canal west of industrial canal crossover road 1E. (pumping lagoon)	X-007	900209-09 A	Beige, fibrous, friable	S, H	Chrysotile Crocidolite	40-50 >1-5	Cellulose 25-35%	Binder Total %:5-10
Top of S. bank (west end) pumping lagoon & on bank (by pump house)	X-008	900209-10 A	Two Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable	1) P, H 2) S, H	Chrysotile Chrysotile Amosite	-	1) Cellulose 1-5% 2) Cellulose 10-20%	1) Binder, quartz Total %:75-85 2) Binder Total %:10-20
West bank pumping lagoon-midway down side-75' from SWcorner, gravelly bank- concrete dump area	X-009	900209-11 A	Three Components: 1) Grey, fibrous, non-friable 2) Cream, fibrous, friable 3) Black, fibrous, sticky	1) P, H 2) S, H 3) S, H	•	2) 60-70	 Cellulose TR None Detected Cellulose 25-35% 	1) Binder, quartz Total%:90-100 2) Binder Total %:20-30 3) Binder Total %:15-25
Parking lot west of pumping lagoon and south (southwest) bank borrow lagoon		900209-12A	Three Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable 3) Black, fibrous, sticky		1) Chrysotile Amosite Crocidolite 2) Chrysotile 3) Chrysotile	5-10 5-10 2) 30-40	 Wollastonite 1-5% Cellulose 10-20% None Detected 	1) Binder, quartz Total %:55-65 2) Binder Total %:30-40 3) Binder Total %:20-30

'Avialyst's Signature Australian Sumples - Surger Page 3 of 5

Client: Illinois	Environmental	Reference:		orms				
	tion Agency		Sample	Sample	Asbestos		Non-asbestos	Non-fibrous
Client I.D.	Client Code	RAI #	Description	Treatmen	I Iype	<u>%</u>	<u>Type</u>	Components
W. of MW1- before hit RR ditch-grassy area	X-011	900209-13A	Two Components: 1) Grey, fibrous, non-friable 2) Black, fibrous, friable	1) P, H 2) S, H			1) Cellulose 10-20% 2) Cellulose 20-30%	1) Binder Total %:40-50 2) Binder, quartz Total %:55-65
Hole-parking area-west of settling basin-8" hole dug under	X-012	900209-14A	Two Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable	1) P, H 2) S, H	, ,	•	1) None Detected 2) None Detected	1) Binder Total %:50-60 2) Binder Total %:40-50

Analyst's Signature Auch Sumfler Hange 4 of 5

NOTE:

Analysis of friable materials performed in accordance with "Interim Method of the Determination of Asbestos in Bulk Samples" by the U.S.E.P.A., 40 CFR Part 763, Subpart F, Appendix A, October 30, 1987. CAUTION: Airborne Asbestos fibers should be strictly avoided as should any activity which may cause the release of asbestos fibers from asbestos-containing materials. Percentages are analyst's best estimate.

This report must not be reproduced except in full and with the approval of the laboratory. This report relates only to the items tested.

H=Homogenize S=Shred N.F.=Non-fibrous Sb=Substantial Tr=Trace D=Dried A=Ashed KEY: P=Pulverize SE=Solvent Extraction

* The U.S.E.P.A. does not indicate the suitability of the aforementioned method for analysis of non-friable bulk samples. As such, the laboratory performs analysis of samples designated as "non-friable" utilizing alternate protocol as specified in the the laboratory Quality Assurance Manual.

Report Approved by

Kirk E. Sweetland, Vice President

Client Reference: Illinois Environmental Protection Agency Johns Manville, Maywood

RAI Job No.: 2-0366-004.01

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this case, the record shall also include information on the location of these naterials.

(2) No provision of this subpart applies to any school if:

(i) The local education agency has onducted abatement programs that esult in the elimination of all friable asbestos materials from the school either by removal or encapsulation of the materials.

(ii) No part of the school building was built before January 1979.

119 References.

(a) General. The following reference contains detailed information on sampling and analysis of friable materials and provides a background on which his part is based. Copies may be obtained from the Document Control Officer, Management Support Division (TF 793), Office of Pesticides and

Substances, Environmental Procuron Agency, Room E-106, 401 M Street SW, Washington, D.C. 20460.

materials in School Buildings: A uidance Document" Part 1. (EPA no. C00090).

PENDIX A TO SUBPART F—INTERIM METHOD OF THE DETERMINATION OF STORE IN BULK INSULATION SAM-

SECTION 1, POLARIZED LIGHT MICROSCOPY

1.1 Principle and Applicability

ulk samples of building materials taken asbestos identification are first examined for homogeneity and preliminary fiber identification at low magnification. Positive identification of suspect fibers is made by llysis of subsamples with the polarized it microscope.

The principles of optical mineralogy are well established. A light microscope ipped with two polarizing filters is used ibserve specific optical characteristics of imple. The use of plane polarized light anows the determination of refractive indices along specific crystallographic axes. I phology and color are also observed. A redation plate is placed in the polarized in the po

ssed polars) allows observation of the

Pt. 763, Subpt. F, App. A

birefringence and extinction characteristics of anisotropic particles.

Quantitative analysis involves the use of point counting. Point counting is a standard technique in petrography for determining the relative areas occupied by separate minerals in thin sections of rock. Background information on the use of point counting and the interpretation of point count data is available.

This method is applicable to all bulk samples of friable insulation materials submitted for identification and quantitation of asbestos components.

1.2 Range

The point counting method may be used for analysis of samples containing from 0 to 100 percent asbestos. The upper detection limit is 100 percent. The lower detection limit is less than 1 percent.

1.3 Interferences

Fibrous organic and inorganic constituents of bulk samples may interfere with the identification and quantitation of the asbestos mineral content. Spray-on binder materials may coat fibers and affect color or obscure optical characteristics to the extent of masking fiber identity. Fine particles of other materials may also adhere to fibers to an extent sufficient to cause confusion in identification. Procedures that may be used for the removal of interferences are presented in Section 1.7.2.2.

1.4 Precision and Accuracy

Adequate data for measuring the accuracy and precision of the method for samples with various matrices are not currently available: Data obtained for samples containing a single asbestos type in a simple matrix are available in the EPA report Bulk Sample Analysis for Asbestos Content: Evaluation of the Tentative Method.

1.5 Apparatus

1.5.1 Sample Analysis

A low-power binocular microscope, preferably stereoscopic, is used to examine the bulk insulation sample as received.

- Microscope: binocular, 10-45X (approximate).
- · Light Source: incandescent or fluorescent.
- Forceps, Dissecting Needles, and Probes
- · Glassine Paper or Clean Glass Plate

Compound microscope requirements: A polarized light microscope complete with polarizer, analyzer, port for wave retardation plate. 360° graduated rotating stage, substage condenser, lamp, and lamp iris.

Polarized Light Microscope: described above

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- Objective Lenses: 10X, 20X, and 40X or near equivalent.
- Dispersion Staining Objective Lens (optional)
- · Ocular Lens: 10X minimum.
- Eyepiece Reticle: cross hair or 25 point Chalkley Point Array.
- Compensator Plate: 550 millimicron retardation.

1.5.2 Sample Preparation

Sample preparation apparatus requirements will depend upon the type of insulation sample under consideration. Various physical and/or chemical means may be employed for an adequate sample assessment.

- Ventitated Hood or negative pressure glove box.
- · Microscope Slides
- · Coversitps
- Mortar and Pestle: agate or porcelain. (optional)
- . Wylie Mill (optional)
- Beakers and Assorted Glassware (optional)
- · Certrifuge (optional)
- · Filtration apparatus (optional)
- · Low temperature asher (optional)

1.6 Reagents

1.6.1 Sample Preparation

- Distilled Water (optional)
- Dilute CH₂COOH: ACS reagent grade (optional)
- Dilute HCL: ACS reagent grade (optional)
- Sodium metaphosphate (NaPO₁)₆ (optional)

1.6.2 Analytical Reagents

Refractive Index Liquids: 1.490-1.570, 1.590-1.720 in increments of 0.002 or 0.004.

- Refractive Index Liquids for Dispersion Staining: high-dispersion series, 1.550, 1.605, 1.630 (optional).
- UICC Asbestos Reference Sample Set: Available from: UICC MRC Pneumoconfosis Unit, Liandough Hospital, Penarth, Glamorgan CF6 1XW, UK, and commercial distributors.
- Tremolite-asbestos (source to be determined)
- Actinolite-asbestos (source to be determined)

1.7 Procedures

Note: Exposure to airborne asbestos fibers is a health hazard. Bulk samples submitted for analysis are usually friable and may release fibers during handling or matrix reduction steps. All sample and slide preparations should be carried out in a ventilated hood or glove box with continuous airflow (negative pressure). Handling of samples without these precautions may result in ex-

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posure of the analyst and contamination of samples by airborne fibers.

1.7.1 Sampling

Samples for analysis of asbestos content shall be taken in the manner prescribed in Reference 5 and information on design of sampling and analysis programs may be found in Reference 6. If there are any questions about the representative nature of the sample, another sample should be requested before proceeding with the analysis.

1.7.2 Analysis

1.7.2.1 Gross Examination

Bulk samples of building materials taken for the identification and quantitation of as bestos are first examined for homogeneity at low magnification with the aid of a ster comicroscope. The core sample may be examined in its container or carefully removed from the container onto a glassine transfer paper or clean glass plate. If possible, note is made of the top and bottom orientation. When discrete strata are identified, each is treated as a separate material so that fibers are first identified and quantified in thal layer only, and then the results for each layer are combined to yield an estimate of asbestos content for the whole sample.

1.7.2.2 Sample Preparation

Bulk materials submitted for asbesto analysis involve a wide variety of matrix materials. Representative subsamples may not be readily obtainable by simple meanin heterogeneous materials, and variousteps may be required to alleviate the difficulties encountered. In most cases, however the best preparation is made by using for ceps to sample at several places from the bulk material. Forcep samples are immersed in a refractive index liquid on a microscopialide, teased apart, covered with a cover glass, and observed with the polarized light microscope.

Alternatively, attempts may be made to homogenize the sample or eliminate interferences before further characterization. The selection of appropriate procedures is dependent upon the samples encountered and personal preference. The following arpresented as possible sample preparation steps.

A mortar and pestle can sometimes bused in the size reduction of soft or loosele bound materials though this may caus matting of some samples. Such samples mabe reduced in a Wylie mill. Apparatus should be clean and extreme care exercised to avoid cross-contamination of sample Periodic checks of the particle sizes should be made during the grinding operation so a to preserve any fiber bundles present in a

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identifiable form. The recommended for sam phibole minerals or ve amphiboles may resul fiber bundles or the pragments with aspec 3:1. Grinding of veroduce fragments wier than 3:1.

Acid treatment ma quired to eliminate i carbonate, gypsum, a are frequently pre: trowelled insulations. be removed by treatn acetic acid. Warm dil may also be used to re rials. If acid treatmen sample at least twice being careful not to during decanting ste filtration of the suspr nificant fiber loss. Th should be 0.45 micror longed acid contact alter the optical chara fibers and should be a

Coatings and bindi to fiber surfaces ma treatment with sod Add 10 mL of 10g/L : solution to a small (0 bulk material in a l tube. For approxima stir the mixture on a an ultrasonic bath an Repeat the series. solids by centrifugat minutes. Wash the : suspending in 10 mL centrifuging. After w pellet in 5 mL distill of the suspension on dry the slide at 110° (

In samples with a k ic or other organic f to ash part of the sai due. Ashing should ' temperature asher. A

Mineral Morph

Chrysotile Wavy fibers (asbesting Splayed in Lorin Aspect rational Serpen Colorless in

Straight rigid

(asbesti-typically torm brown, non-grunerite) so Opaque present

Amosite

Environmental Protection Agency

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identifiable form. These procedures are not recommended for samples that contain amphibole minerals or vermiculite. Grinding of amphiboles may result in the separation of fiber bundles or the production of cleavage fragments with aspect ratios greater than 3:1. Orinding of vermiculite may also produce fragments with aspect ratios greater than 3:1.

Acid treatment may occasionally be required to eliminate interferences. Calcium carbonate, gypsum, and bassanite (plaster) are frequently present in sprayed or trowelled insulations. These materials may

emoved by treatment with warm dilute basels acid. Warm dilute hydrochloric acid may also be used to remove the above materials. If acid treatment is required, wash the sample at least twice with distilled water, being careful not to lose the particulates during decanting steps. Centrifugation or filtration of the suspension will prevent significant fiber loss. The pore size of the filter should be 0.45 micron or less. Caution: prolonged acid contact with the sample may "be the optical characteristics of chrysotlie its and should be avoided.

Coatings and binding materials adhering to fiber surfaces may also be removed by treatment with sodium metaphosphate. Add 10 mL of 10g/L sodium metaphosphate solution to a small (0.1 to 0.5 mL) sample of bulk material in a 15-mL glass centrifuge tube. For approximately 15 seconds each, stir the mixture on a vortex mixer, place in an ultrasonic bath and then shake by hand. Repeat the series. Collect the dispersed

's by centrifugation at 1000 rpm for 5 sutes. Wash the sample three times by suspending in 10 mL distilled water and recentrifuging. After washing, resuspend the pellet in 5 mL distilled water, place a drop of the suspension on a microscope slide, and dry the slide at 110' C.

In samples with a large portion of cellulosic or other organic fibers, it may be useful to ash part of the sample and view the residue. Ashing should be performed in a low temperature asher. Ashing may also be per-

formed in a muffle furnace at temperatures of 500° C or lower. Temperatures of 550° C or higher will cause dehydroxylation of the asbestos minerals, resulting in changes of the refractive index and other key parameters. If a muffle furnace is to be used, the furnace thermostat should be checked and calibrated to ensure that samples will not be heated at temperatures greater than 550° C.

Ashing and acid treatment of samples should not be used as standard procedures. In order to monitor possible changes in fiber characteristics, the material should be viewed microscopically before and after any sample preparation procedure. Use of these procedures on samples to be used for quantitation requires a correction for percent weight loss.

1.7.2.3 Fiber Identification

Positive identification of asbestos requires the determination of the following optical properties.

- Morphology
- Color and pleochroism
- Refractive indices
- · Birefringence
- Extinction characteristics
- · Sign of elongation

Table 1-1 lists the above properties for commercial asbestos fibers. Figure 1-1 presents a flow diagram of the examination procedure. Natural variations in the conditions under which deposits of asbestiform minerals are formed will occasionally produce exceptions to the published values and differences from the UICC standards. The sign of elongation is determined by use of the compensator plate and crossed polars. Refractive indices may be determined by the Becke line test. Alternatively, dispersion staining may be used. Inexperienced operators may find that the dispersion staining technique is more easily learned, and should consult Reference 9 for guidance. Central stop dispersion staining colors are presented in Table 1-2. Available high-dispersion (HD) Houlds should be used.

TABLE 1-1-OPTICAL PROPERTIES OF ASBESTOC FIBERS

Mineral	}	Retractive	r indices*	Birefrin-		Sign of elonation (length slow)
	Morphology, color*	a	γ	gence	Extinction	
Chrysotile (asbesti- form serpen- tine)	Wavy libers Fiber bundles have solayed ends and likinks. Aspect ratio typically > 10.1 Colorless i nonpleochroic	1 493-1 560	1 517- 1 562' Inormally 1 5561	008	to liber length	
Amosite (asbesti- form rundrite).	Straight, rigid fibers. Aspect ratio typically >10.1. Colorless to brown, nonpleochroic or weakly sc. Opaque inclusions may be present.	1 635-1 696	1 655- 1 729' Ingrmally 1 696- 1 710	020- 033	ito fiber length	(length slow)

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TABLE 1-1—OPTICAL PROPERTIES OF ASBESTOC FIBERS—Continued

Mineral		Refractive	ndices*	Breton-		Sign of
	Morphology, color*	α	γ	gence	Extinction	elonation
Crocidolite (asbesti- form Riebeck- ite).	Straight, rigid libers. Thick fibers and bundles common, blue to purple-blue in color. Pleochroic Birefringence is generally masked by blue color.	1.654-1.701	1.668- 1.717 * (normally close to 1.700).	.014~.016	No fiber length	(length fasi)
Anthophyl- ide- asbestos.	Straight hibers and accular cleav- age fragments ^e Some composite libers. Aspect ratio < 10:1. Color- less to light brown.	1.596-1.652	1 615- 1.6761	.019024	Itlo fiber length.	(length slow)
Tremokte- actinolite- asbestos.	Normally present as accoular or prematic cleavage fragments. Single crystals predominate, aspect ratio < 10:1. Colorless to pale green.	1.599-1.668	1.622- 1.608'.	023020	Oblique extinction, 10-20 for fragments Composite libers show # extinction.	(length slow)

*From reference 5; colors cited are seen by observation with plane polarized light.
*From references 5 and 8.
*From references 5 and 8.
*From references 5 and 8.

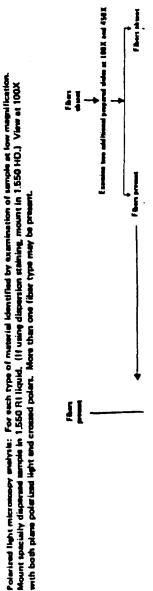
*From references 5 and 8.

*From references 5 and 8.

*From references 5 and 8.

*Literal from references 5 and

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Polarized light microscopy Mount spacially dispersed with both plans polarized

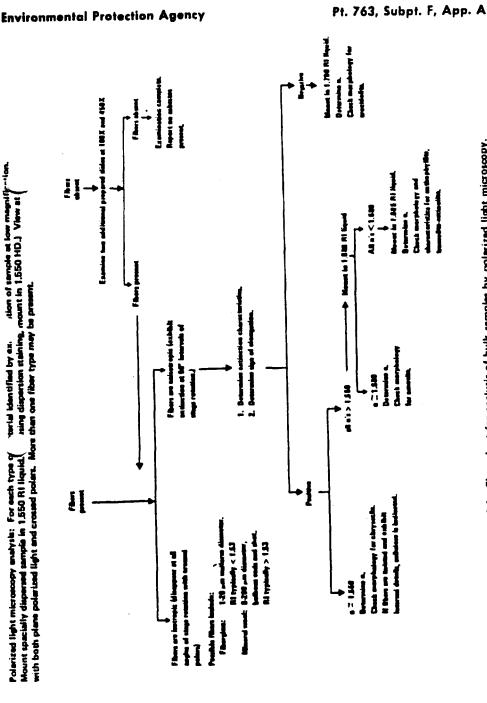


Figure 1-1. Flow chart for analysis of bulk samples by polarized light microscopy.



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TABLE 1-2—CENTRAL STOP DISPERSION STAINING COLORS.

Mineral	RI Liquid	נק	ווֹר
Chrysotile	1 550***	Blue	Blue-magenta
Amosile	1.680	Blue-magenta to pale blue	Golden-yellow
	1.550***	Yellow to white	Yellow to white
Crocidokte*	1.700	Red magenta	Blue-magenta
	1.550***	Yellow to white.	Yellow to white
Anthophylide	1.605 ^{mil}	Blue	Gold to gold- magents
Tremokte	1 605****	Pale blue	Gold
Actinoiste	1.605***	Gold-magenta to blue.	Gold
	1.630****	Magenta	Golden-yellow

^{*}From reference 9

Asbestos quantitation is performed by a point-counting procedure or an equivalent estimation method. An ocular reticle (crosshair or point array) is used to visually superimpose a point or points on the microscope field of view. Record the number of points positioned directly above each kind of particle or fiber of interest. Score only points directly over asbestos fibers or nonasbestos matrix material. Do not score empty points for the closest particle. If an asbestos fiber and a matrix particle overlap so that a point is superimposed on their visual intersection, a point is scored for both categories. Point counting provides a determination of the area percent asbestos. Reliable conversion of area percent to percent of dry weight is not currently feasible unless the specific gravities and relative volumes of the materials are known

For the purpose of this method, "asbestos fibers" are defined as having an aspect ratio greater than 3:1 and being positively identified as one of the minerals in Table 1-1.

A total of 400 points superimposed on either arbestos fibers or nonasbestos matrix material must be counted over at least eight different preparations of representative subsamples. Take eight forcep samples and mount each separately with the appropriate refractive index liquid. The preparation should not be heavily loaded. The sample should be uniformly dispersed to avoid overlapping particles and allow 25-50 percent empty area within the fields of view. Count 50 nonempty points on each preparation, using either

- A cross-hair reticle and mechanical stage; or
- A reticle with 25 points (Chalkley Point Array) and counting at least 2 randomly selected fields.

For samples with mixtures of isotropic and anisotropic materials present, viewing the

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sample with slightly uncrossed polars or the addition of the compensator plate to the polarized light path will allow simultaneous discrimination of both particle types. Quantitation should be performed at 100X or at the lowest magnification of the polarized light microscope that can effectively distinguish the sample components. Confirmation of the quantitation result by a second analyst on some percentage of analyzed samples should be used as standard quality control procedure.

The percent asbestos is calculated as follows:

% asbestos = (a/n) 100%

where

a = number of asbestos counts.

n=number of nonempty points counted (400).

If a=0, report "No asbestos detected." If 0 < a < 3, report "<1% asbestos".

The value reported should be rounded to the nearest percent.

1.8 References

- 1. Paul F. Kerr, Optical Mineralogy, 4th ed., New York, McGraw-Hill, 1977.
- 2. E. M. Chamot and C. W. Mason, Handbook of Chemical Microscopy, Volume One. 3rd ed., New York: John Wiley & Sons, 1958.
- 3. F. Chayes, Petrographic Modal Analysis: An Elementary Statistical Appraisal, New York: John Wiley & Sons, 1956.
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- 5. U.S. Environmental Protection Agency. Asbestos-Containing Materials in School Buildings: A Guidance Document, Parts 1 and 2, EPA/OTS No. C00090, March 1979.
- 6. D. Lucas, T. Hartwell, and A. V. Rao, Asbestos-Containing Materials in School Buildings: Guidance for Asbestos Analytical Programs, EPA 560/13-80-017A, U.S. Environmental Protection Agency, December 1980, 96 pp.
- 7. D. H. Taylor and J. S. Bloom, Hexametaphosphate pretreatment of insulation samples for identification of fibrous constituents, *Microscope*, 28, 1980.
- 8. W. J. Campbell, R. L. Blake, L. L. Brown, E. E. Cather, and J. J. Sjoberg. Sciected Silicate Minerals and Their Asbesti form Varieties: Mineralogical Definitions and Identification-Characterization, U.S. Bureau of Mines Information Circular 8751, 1917.
- 9. Walter C. McCrone, Asbestos Particl-Atlas, Ann Arbor: Ann Arbor Science Publishers, June 1980.

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SECTION 2. X-RAY P

2.1 Principle a

The principle of X-1 (XRD) analysis is woolld, crystalline mat impingent beam of pa X-rays whenever Brag $\lambda = 2d \sin \theta$.

is satisfied for a part the crystal lattice, who λ = the X-ray waveler.

- d = the A-ray waveler.

 d = the interplanar s;

 flecting lattice pla
- the angle of inciray beam and t planes.

By appropriate orient tive to the incident 2 tion pattern can be ge cases, will be uniquely the chemical composithe crystalline phases

Unlike optical mether, XRD cannot deter ogy. Therefore, in as does not distinguish nonfibrous forms of t phibole minerals (I when used in conjuncthods such as pola (PLM), XRD techniquable analytical metherion and characteriz minerals in bulk mate.

TABLE 2-2

Minerals
Chrysotile
"Amoste"
Anthophylide
Anthophyline

*This information is intended be referred to, to ensure com amosite, crocidolite, tremolite, *Fibrosity questionable

Accurate quantitati in bulk samples by XF ent on particle size d size, preferred orient sorption effects, and c ard reference and si

^{*}Blue absorption color.

^{1.7.2.4} Quantitation of Asbestos Content

5.2.2.2 Carefully determine diameter (d) of Walton-Beckett graticule, use the following equation to determine MFA:

$$MFA = \pi (d/2)^2$$

5.2.3 Determining the Phase-Shift Detection Limit of Microscope

To be performed initially upon receipt of microscope and periodically thereafter. Procedure is as follows:

- 5.2.3.1 With 40X objective in place, focus sharply on the blocks of grooved lines on an HSE/NPL phase contrast test slide.
- In order for the microscope to be used for fiber counting the microscope optics must be capable of resolving the three most visible blocks of grooves clearly, the fourth and fifth most visible blocks at least partially, and the sixth and seventh blocks of grooves must be totally invisible. Failure may indicate that microscope has too high or low a resolution to be used for fiber counting.
- 5.2.3.3 Deterioration of resolution of a microscope may indicate need for cleaning and/or service of microscope

5.2.4 Determining Resolution of Analysts' Eyes

To be performed by each analyst at least annually.

- 5.2.4.1 Once microscope resolution has been established through the process described in 5.2.3, the analyst is asked to determine the number of blocks of grooved lines that they see. The analyst must make certain to adjust interocular distance and ocular focus for their use.
- 5.2.4.2 If analyst is unable to discern the fifth block of grooves at least partially, then the analyst is withheld from performing fiber-counting. Retesting will be permitted.

5.3.0 Polarized Light Microscope (PLM)

5.3.1 Alianment and Set-up

5.3.1.1 <u>Initial</u>

Calibration + Alignment from our Quality Assurance Manual

To be performed at the beginning of each work shift. It is assumed that analyst will adjust interpupillary distance and ocular focus to their best advantage.

- 5.3.1.1.1 With illumination on, analyzer out, and dispersion staining objective in line, focus on any prepared slide. Note: Be certain that no stop is in line of focus. Check with Bertrand lens.
- 5.3.1.1.2 Check centration of rotating stage and adjust with stage centering screws.

- 5.3.1.1.3 Close field and aperture diaphragms and rack condenser to bring field diaphragm into sharpest possible focus.
- 5.3.1.1.4 Center field diaphragm image with condenser centering screws. Open field diaphragm to just fill field of view with light.
- 5.3.1.1.5 Center other objectives using objective centering screws on rotating nosepiece.

Note: During use of microscope to examine samples, each time a different objective is rotated into place, field daphragm should be closed, focused and opened to just fill field of view. This should also be performed each time a different slide is examined. Objective centering should be corrected whenever it is observed to have changed.

5.3.1.2 Central Stop Alignment

To be performed following initial alignment described in 5.3.1.1 and before the beginning of each work shift.

- 5.3.1.2.1 With dispersion staining objective in line, insert Bertrand lens.
- 5.3.1.2.2 Bring largest central stop into field of view.
- 5.3.1.2.3 Center central stop so that as aperture diaphragm is closed, the image of the aperture diaphragm contracts behind the stop in a concentric fashion.
- 5.3.1.2.4 Remove Bertrand lens.

5.3.1.3 Polarizer Alignment

To be performed when new microscope is put into service or following moving of microscope. To be checked and adjusted periodically as needed.

- 5.3.1.3.1 Begin with red dots on intermediate attachment and microscope stand aligned. Bring 10X objective into light path.
- 5.3.1.3.2 Set both polarizer and analyzer at position "0" to attain "crossed filter" position. Insert analyzer.
- 5.3.1.3.3 Place orientation plate on the center of the stage.
- 5.3.1.3.4 Focus on edge of plate and rotate stage until it most darkens. Move orientation plate so that edge of plate nears x-axis cross line.
- 5.3.1.3.5 Withdraw analyzer.
- 5.3.1.3.6 Loosen observation tube clamping screw and rotate observation tube so that x-axis cross line is parallel to edge of plate. Retighten observation tube clamping screw.
- 5.3.1.3.7 Replace orientation plate with specimen slide, selecting a transparent area.

- 5.3.1.3.8 Insert analyzer.
- 5.3.1.3.9 Loosen polarizer clamping screw, rotate polarizer until darkest field is obtained. Tighten polarizer clamping screw.

5.3.2 Cross Line Reticle Calibration

To be performed when new microscope is put into service and/or when reticle or other optical elements are changed.

- 5.3.2.1 Place stage micrometer in center of stage. With 10X objective in light path and analyzer withdrawn, focus sharply on micrometer.
- 5.3.2.2 Line up micrometer with x-axis line of cross line reticle and carefully measure length of scale on reticle. Record measurement specifically for that objective.
- 5.3.2.3 Repeat measurement of reticle scale and recording of measurement for each objective on microscope.
- 5.3.2.4 Calculate length of reticle units for each objective using following formula:

of micrometers measured = micrometers / crossline reticle unit crossline reticle units measured

Note: Not all PLMs in each lab will be equipped with a scaled crossline reticle. At least one PLM per lab is to have this accessory.

5.4 Adjustments to Optical Microscope for Individual Analyst

Each analyst should routinely make these adjustments prior to use of a scope. Rechecking adjustments hourly is highly recommended to reduce eye strain.

- 5.4.1 Adjust interpupillary distance of oculars simply by pulling apart or pushing together by the knurled dovetail slides of the left and right eyepiece tubes. All microscopes used by RAI automatically compensate for changes in focal length that might be associated with this process.
- 5.4.2 Ocular focus and adjustments are performed by the following:
- 5.4.2.1 With interpupillary distance adjusted for analyst, cover the left eye and look into right eyepiece with right eye.
- 5.4.2.2 Focus the reticle by rotating the top lens of the ocular. Note: Reticle eyepieces are to always be inserted in right ocular tube.
- 5.4.2.3 With left eye still covered, focus sharply on very small object on a specimen slide with coarse and fine microscope focus knobs.

5.4.2.4 Uncover left eye, cover right eye. Using Independent focusing adjustment at the base of the ocular, focus sharply on source particle as before for the left eye/left ocular combination.

All calibrations / verifications are to be documented in the daily Calibration Record form in each analyst's notebook.

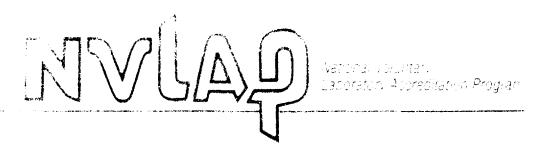
5.5. Verification of the Refractive Index of Immersion Oils

The refractive index of all oils used routinely in the central stop dispersion staining method for PLM shall be verified semi-annually and/or when evidence suggests poor or deteriorating performance of the oil. Verification shall be performed through the use of a refractometer which has been calibrated and is capable of maintaining a constant temperature of 25 degrees C. At least one bottle of each commonly used oil (High Dispersion oils 1.55, 1.605 and 1.68) from each lot received shall be subjected to this verification procedure. In addition, RAI will maintain a set of oils of refractive indices 1.400-1.780 at increments of 0.004 which shall be calibrated/ verified on an annual basis.

5.6 Maintenance and Monitoring of HEPA Filter Hoods in the PLM Laboratory

RAI will provide Class I Biohazard HEPA-filter hoods for each bulk analysis workstation. Optimal flow rates will be verified through the use of a smoke test which shall be performed at a frequency of once per month. Each analyst will report any perceived deterioration of performance of their hood immediately to the Laboratory Operations Manager who will methodically isolate the source of the problem. In addition, each HEPA filter shall be routinely inspected for gross contamination. It is assumed that the HEPA filter is in good operating condition if the hood is capable of maintaining the inward velocity indicated in the manufacturer's specifications.

In the event that a HEPA filter becomes clogged or damaged and requires service, it shall be treated as an asbestos-containing material and be handled with the utmost care. The individual disassembling the unit shall wear a fitted respirator, double-bag the contaminated filter and dispose of it in a manner appropriate for asbestos waste.



SCOPE OF ACCREDITATION

ASBESTOS FIBER ANALYSIS

NVLAP LAB CODE 1408 00

Randolph & Associates, Inc. 8901 N. Industrial Rd. Peoria, IL 61615

Kirk Sweetland

Phone: 309-692-4160

Accreditation Renewal Date: April 1, 1990

NVLAP Test

Method Code

Test Method Designation

18/A01

40 Code of Federal Regulations Chapter I (1-1-87 edition) Part 763, Subpart F, Appendix A, pages 293-299 or the current U.S. Environmental Protection Agency method for the analysis of asbestos in building materials by polarized light microscopy.



the National Institute of Standards and Tachne dow

United States Department of Commerce National Institute of Standards and Technology



RANDOLPH & ASSOCIATES, INC. PEORIA, IL

is recognized under the National Voluntary Laboratory Accreditation Program
for satisfactory compliance with criteria established in Title 15. Part 7 Code of Vederal Regulations
Accreditation is awarded for specific services, listed on the Scope of Accreditation, for:

ASBESTOS FIBER ANALYSIS

April 1, 1990

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For the National Institute of Standards and Technologic

Copy



Randolph & Associates, Inc.

2201 W. Townline Road, Peoria, Illinois 61615 FAX 692-9364 Telephone 309-692-4160

Client:

Illinois Environmental Protection Agency

Division of Land Pollution Control

2200 Churchill Road

Springfield, Illinois 62706

Contact:

Kurt Neibergall, Project Manager

Analysis: Routine/Unit Cost

ASBESTOS BULK SAMPLE EVALUATION

POLARIZED LIGHT MICROSCOPY (PLM) TECHNIQUE

Date Analyzed:

2/9/90

Date Received:

2/9/90

Client Reference: Johns Manville

Maywood

Client P.O.#:

RAI Job #:

2.0366.004.01

Type of Sampl	e: Soil/Debr	is Samples	Sampled b	y: Client				Analyst:	Anita	Sumpter-Sturgies
	· · · · · · · · · · · · · · · · · · ·				 F	lbre	ous F	orms		
Client I.D.	Client Code	RAL #	Sample <u>Description</u>	Sample <u>Treatment</u>	Asbestos Type		<u>%</u>	Non-asbestos <u>Type</u>		Non-fibrous Components
Surface grab sample 50' east of NE corner, misc. pit (15' from east edge of roadway)	X-001	900209-03 A	Two Components: 1) Brown, fibrous, non-friable 2) Grey, fibrous, friab	1) P, H 2) S, H le	•	•		1) None Detected 2) Cellulose 1-5%		1) Binder, quartz Total %:55-65 2) Binder, quartz Total %:10-20
East edge of road-80 yd. north of NE corner misc. pit (20' west of MW #7)	X-002	9002 09-04A	Two Components: 1) Black and grey, fibrous, non-friable 2) Grey and brown, fibrous, friable	1) P, H 2) S, H	Chrysotile Chrysotile			1) Cellulose TR 2) Cellulose TR		1) Binder Total %:50-60 2) Binder, quartz Total %:85-95

Cllent: Illinois Environmental	Reference:	Johns Manville			F	ibr	ous F	orms	
Protection Agency		Sample	Sample		Asbestos		٥/	Non-asbestos <u>Type</u>	Non-fibrous Components
Client I.D. Client Code	RAI #	<u>Description</u>	Treatmen		Type	4.	<u>%</u>		•
15' East of X-003 roadway-pipe/ siding debris 70 yds. from bend in road to west, -surface grab sample	900209- 05A	Two Components: 1) Brown, fibrous, non-friable 2) Grey and brown, fibrous, friable	1) P, H 2) S, H		Chrysotile Chrysotile Amosite			1) Wollastonite 1-5% 2) Cellulose 1-5%	1) Binder, quartz Total %:45-55 2) Binder, quartz Total %:45-55
East edge of X-004 roadway-sludge / pipe-gravelly 30' north of MW5	900209-06 A	Two Components: 1) Black, fibrous, non-friable 2) Grey, fibrous, friable	1) P, H 2) S, H					1) None Detected 2) None Detected	1) Binder Total %:20-30 2) Binder Total %:10-20
Undercut-bank- X-005 indutrial canal (south side) approx. 20 ydwest of new overflow structure	900209-07 A	Grey, fibrous, friable	S, H	CI	nrysotile	50	-60	Cellulose 5-10%	Binder Total %:20-30
1/2 way down X-006 west bank industrial canal -50" from NW corner (surface grab)	900209-08 A	Three Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable 3) Black, fibrous, gelatinous	1) P. H 2) S, H 3) S, H	2)	Chrysotile Chrysotile Crocidolite None etected	2)		1) Cellulose TR 2) None Detected 3) Cellulose 60-70%	1) Binder Total %:85-95 2) Binder Total %:10-20 3) Binder Total %:20-30

Analyst's Signature Justa Jumplus Jungies Page 2 of 5

CIIent: Illinois	Environmental	Reference:	Johns Manville		F	orms		
Protect <u>Client I.D.</u>	ion Agency Cllent Code	RAI #	Sample <u>Description</u>	Sample Treatmen	Asbestos it Type	<u>%</u>	Non-asbestos <u>Type</u>	Non-fibrous <u>Components</u>
Southeast corner-canal west of industrial canal crossover road 1E. (pumping lagoon)	X-007	900209- 09A	Beige, fibrous, friable	S, H	Chrysotile Crocidolite	40-50 >1-5	Cellulose 25-35%	Binder Total %:5-10
Top of S. bank (west end) pumping lagoon & on bank (by pump house)	X-008	900209-10 A	Two Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable	1) P, H 2) S, H	 Chrysotile Chrysotile Amosite 		1) Cellulose 1-5% 2) Cellulose 10-20%	1) Binder, quartz Total %:75-85 2) Binder Total %:10-20
West bank pumping lagoon-midway down side-75' from SWcorner, gravelly bank- concrete dump area	X-009	900209-11 A	Three Components: 1) Grey, fibrous, non-friable 2) Cream, fibrous, friable 3) Black, fibrous, sticky	1) P, H 2) S, H 3) S, H		2) 60-70	1) Cellulose TR 2) None Detected 3) Cellulose 25-35%	1) Binder, quartz Total%:90-100 2) Binder Total %:20-30 3) Binder Total %:15-25
Parking lot west of pumping lagoon and south (southwest) bank borrow lagoon		900209-12 A	Three Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable 3) Black, fibrous, sticky		-	5-10 5-10 2) 30-40	 Wollastonite 1-5% Cellulose 10-20% None Detected 	1) Binder, quartz Total %:55-65 2) Binder Total %:30-40 3) Binder Total %:20-30

Analyst's Signature Austa Sumplus Surger Page 3 of 5

Cilent: Illinois	Environmental	Reference:	Johns Manville	Fibrous Forms			orms	
Protec	tion Agency		Sample	Sample	Asbestos		Non-asbestos	Non-fibrous
Client I.D.	Cllent Code	RAL #	<u>Description</u>	<u>Treatmen</u>	<u>t Type</u>	<u>%</u>	<u>Type</u>	Components
W. of MW1- before hit RR ditch-grassy area	X-011	900209-13A	Two Components: 1) Grey, fibrous, non-friable 2) Black, fibrous, friable	1) P, H 2) S, H	Chrysotile Chrysotile	•	1) Cellulose 10-20% 2) Cellulose 20-30%	1) Binder Total %:40-50 2) Binder, quartz Total %:55-65
Hole-parking area-west of settling basin-8" hole dug under gravel	X-012	900209-14 A	Two Components: 1) Grey, fibrous, non-friable 2) Grey, fibrous, friable	1) P, H 2) S, H	•	•	None Detected None Detected	1) Binder Total %:50-60 2) Binder Total %:40-50

Analyst's Signature Auch Sunfty Page 4 of 5

NOTE:

Analysis of friable materials performed in accordance with "Interim Method of the Determination of Asbestos in Bulk Samples" by the U.S.E.P.A., 40 CFR Part 763, Subpart F, Appendix A, October 30, 1987. CAUTION: Airborne Asbestos fibers should be strictly avoided as should any activity which may cause the release of asbestos fibers from asbestos-containing materials. Percentages are analyst's best estimate.

This report must not be reproduced except in full and with the approval of the laboratory. This report relates only to the items tested.

H=Homogenize S=Shred N.F.=Non-fibrous Sb=Substantial Tr=Trace D=Dried A=Ashed KEY: P=Pulverize SE=Solvent Extraction

* T' U.S.E.P.A. does not indicate the suitability of the aforementioned method for analysis of non-friable bulk samples. As such, the laboratory performs analysis of samples designated as "non-friable" utilizing alternate protocol as specified in the the laboratory Quality Assurance Manual.

Report Approved by

Kirk E. Sweetland, Vice President

Client Reference: Illinois Environmental Protection Agency

Johns Manville, Maywood

RAI Job No.: 2.1602.001.01

page 5 of 5

LEPA CHAIN & CONTROL			Seal #F, TAPE led: 2/6/90 By: AWW
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Seal #E-TAPE Date Sealed: 2690 By: AWW Facil: Site Inventory # :0971900014 Site Billing edd: 12-52195 Name Project Hanager : KURT NEIBERGALL SAMPLE DATE SAMPLE I.D. SAMELE TIME 3-00 PM X-010 Collector Comments: ANTHOLOT WEST OF PWANTS LABORY SUNGE & ATTINGENT SUNGER & ATTINGENT & ATTINGENT SUNGER & ATTINGENT & ATTING Collector Comments: APPLY to LOT WEST OF PUMPING LABOUR AND SAME BORDE READ Sampler Signature : Division/Company INA DUR RANS FILTERING Blo(Y/N) Date | Time KULT HEUBELGAU STEWEHARRUB CUSTODY CHRONICLE incertify that I received the sample shipping container with the shipping container sealed and intact Opened by (print):_________Signature:_____ Da':: _____ Time: ____ Seal #: ____ IntactF: Y / N i certify that the sample listed above was collected and placed in bottles in my presence, that each bottle was placed intect in the famile shipping container and that I sealed the sample shipping container at the date and time listed below Sealed by (print):______ Signature: Courier - sample pickup; ______

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Facility Name 1000 FANVILLE		Site Inventory #		
Region Participal		Site Billing Code Project Manager	:LP-52195	
County				
	SAMPLE I.D	SAMPLE DATE 2-7-90		1-55PM
Sample Appearance :C	X-00H	> FIBLOUS		1-3211
Sample Appearance :	HMKY SIND	SE-WHITE/BLUEIS	4 GREY -	- BADWA SOIL MATRY
Collector Comments: E. E.	DOK OF KOADY			
Sampler Signature :	XI) THE	rision/Compan	y IRPA	OURC/RAYS
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in certify that I received the sample ship	ping container with the	shipping contained sealed and i	rtact	
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IEPA - CLP CHAIN OF CUSTODY		Date Seale	Seal #: <u>E.</u> d: 2/6/90	TAPE By: AWW
Facility The Name of the Pacific Control of t		Site Inventory		
Regio Counti		Site Billing Co Project Manager	de: LP-52195	RGALL
	SAMPLE I D	SAMFLE	DATE S/	MELF TIME
	X-003	2-7-90) =	50 PM
Sample Appearance :	SOLL - DARK BRO	WM - W/ SLUDGE	AM 300M MAN	PERIALS I
Collector Comments: $\frac{15}{2}$	EAST OF ROADWAY	PIPE SIDHE DEBRIS FROM BEHD IN ROAD Division/Comp	SAMPLE SULFACE TO LIESTEPA/DU any	CHUNKY MLS-SOL-SANDT CRAUS WILLIAMS WI
	A STATE OF THE PARTY OF THE PAR	NALYSIS		RING HEW
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			2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	
KUKT NEIBERGALI I certify that I received the sample		FODY CHRONICLE	1 I	
· ·		•		
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I certify that the sample listed about sample shipping container and that I		• •		sed infact in the
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Date:T	ime:	Seal #:		
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LEPA - GE CFAIN T- COLT		Date S	Seal	#:E-TA	PE NW
Facilia Name Rege VIII County		Site Invent Site Billin Project Man	ory # :0971 g Code: <u>LP-</u> 9	1900014	
Samp le Appearance : 1	SAMPLE I.E X-002	2-	OF NE COS	90MFL 9 1:40	PM
Collector Comments: Collector Comments: Collector Comments: Collector Comments: Campler Signature :	CRAVELLY KAN	DY DARK BROWN	UM- MIXE -Moist (1) Company ID	DW/WHITE	CRUMBLY - SCRAPES M SHARDANI
	r. sistos	ANALYSIS	I C	FILTERIN Y(N): Date A/:	the state of the s
KURT NEIBERGALL I certify that I received the sample	CHAIN OF COS	ETODY CHRONICL			<u>-</u>
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I certify that the sample listed above sample shipping container and that I					avitoria di
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EPA - N.P.		- 14 - 12	\nku 81	• 4	E. TA1 19084:	
acilii ame egi		Site I	ate Sealed inventory # Silling Cod it Manager	:0971900	014 5.	
	SAMPLE X - OC	E I.D.	2-7-90		1:25 P	
ample Appearance	: MED BROWN	- SAMDY SO	11MOIST	- SOME	GRAUEL	
ollector Comments	: SURFACE GRA	B, SAMPLE	50 E. of	HE COPH	er misc	
empler Signature	- Kuth JA	H-AC	sion/Compa	ny tery	Durger	777
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KWRT D. HEIBSTEAN	L STEVE DAVIS	> F CUSTODY CHE	RONICLE			
entify that I received the s	ample shipping container	with the shipping co	ntainer sealed and	intact		
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ertify that the sample liste sole shipping container and t	d above was collected and	d placed in bottles in	n my presence, the	t erch bilt e .		
ealed by (print):		Sign	nature:	. In the case of t		
ate:	Time:	Seal #	<u> </u>			
ourier - sample p	ickup:				gar gangan make garap akaka sakka sasar	
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SAMPLING MAP OF 2-7-90 BY BRAD BRADLEY (US. EPA) E. MEYERS (EDI)
KURT NEIBERGAUL (I EPA) AND STEVE DAVIS (IEPA) SAM JUNG (C.R.A.)
MANVILLE

